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Descripti n

The present invention relates to base oil composition which have relatively lower toxicity than oils having relatively higher aromatic hydrocarbon content for use especially in drilling fluid formulations.

Drilling fluids generally contain as essential ingredients a base fluid component such as diesel or water, emulsifiers, surfactants or wetting agents, clays, water, chlorides of sodium and calcium, and weighting materials. The base oil component is usually either a water-based, i.e. water forms the continuous phase, or, an oil-based, i.e. oil forms the continuous phase. Normally, water is purposely added to the latter types to give oil based (water-in-oil) emulsions; these are the so-called 'invert emulsions'. The function of the drilling fluid is to (a) cool and lubricate the drill bit and drill string, (b) remove drilled solids, (c) suspend drilled cuttings and other debris by gelation, (d) facilitate control of surface pressures, and (e) prevent caving of formations and to shore up sides of the bore holes.

The type of drilling fluid formulation used will depend amongst others upon the conditions of drilling and the geology of the formation being drilled. In addition the properties of the drilling fluid formulation are also important. These properties include its rheology, density, viscosity, gelation characteristics, ease and rate of important. These properties include its rheology, density, viscosity, gelation characteristics, ease and rate of important. These properties include its rheology, density, viscosity, gelation characteristics, ease and rate of important. The viscosity properties in the conditions invertigate the conditions invertigate the conditions of the conditions of the conditions are preferred.

Hitherto, the base oil component used in these drilling fluids has almost exclusively been diesel. The reason for this is that diesel is relatively inexpensive and acts quickly upon the organophilic clays present as viscosifiers in drilling fluids and accelerates the swelling and gelation of the clay. This is a necessary mechanism to impart optimum rheology to the fluid. However, in spite of its good performance, diesel is undesirable due to the presence of a high concentration (up to 25% w/w) of aromatic hydrocarbons therein, of which up to 2% are polycyclic, which are toxic. Moreover, a portion of the used drilling fluids based on diesel are usually discharged into the sea because they are retained on the rock cuttings which are difficult to clean and thereby risk pollution of the environment.

Efforts have been made to mitigate this problem by using low toxicity oils. However such oils if used alone require prolonged lhearing and/or higher levels of viscosifiers such as clays to achieve the desired initial rheology. It has been recognised that this deficiency of low toxicity oils could be mitigated by using polar activators to make up for the loss of the polar aromatic components in diesel. However, such efforts have been relatively unsuccessful and the drilling industry has become resigned to the added expense of extra shearing or the use of up to 50% more viscosifier clay to replace diesel in drilling fluids.

It has now been found that by choosing a base oil of low toxicity and combining it with an appropriate polar activator, the problems of toxicity, shearing, stability, gelation etc in drilling fluids containing conventional base oils can be mitigated.

Accordingly, the present invention is a base oil composition suitable for use in drilling fluids, said composition comprising an oil component with an aromatic hydrocarbon content of less than 10% w/w and a polar activator which is an ether alcohol or a mixture of ether alcohols.

The oil component in the base oil composition may be suitably kerosine, gas oils low in aromatic hydrocarbons (also known as technical gas oil) or an oil which is derived by diluting diesel with another oil which is low in aromatic hydrocarbon content such that the total aromatic hydrocarbon content of the composition is less than 10% w/w. Such oils are hereafter termed as 'low toxicity oils'.

The polar activator used in the base oil composition of the present invention is suitably a polar ether alcohol i.e. an ether alcohol which has inherent polarity.

The ether alcohols (also known as "glycol ethers") used contain one or more ether linkages. The ether alcohols may be aliphatic, aromatic or mixtures of these. They are suitably derivable from organic compounds having an -OH group (e.g. alcohols or phenols) such as isobutanol or phenol and an alkylene oxide but are preferably derivable from an aromatic compound having -OH groups such as e.g. phenol or a phenoxy propanol and an alkylene oxide which suitably has 2-4 carbon atoms, preferably 2-3 carbon atoms e.g. ethylene oxide or propylene oxide.

Thus the ether alcohol activator can be one or more of a butoxypropanol, isobutoxypropanol, phenoxypropanol, phenoxypropanol, a phenoxypropoxypropanol or substituted derivatives thereof such that the substituents do not adversely affect the performance of the base oil composition when used in drilling fluids.

The relative amounts of the oil component and the polar activator in the base oil composition is suitably such that it contains from 1-5% by weight, preferably from 1.5-3.5% w/w of the activator.

Under certain conditions aromatic ether alcohols such as phenoxypropanol have a tendency to separate out from the base oil. This is particularly so at low temperatures. In such a case it is preferable to use the aromatic ether alcohol in conjunction with an aliphatic ether alcohol such as ethoxypropoxypropanol. For

instance, at base oil temperatures below 15°C, using a mixture of ether alcohols containing from 20-50% w/w of an aliphatic ether alcohol prevents such separation. In addition, the presence of an aliphatic ether alcohol in the mixture depresses the freezing point.

As mentioned previously, drilling fluids contain in addition to the base oil composition conventional components such as clays, water, salts, e.g. chlorides of calcium and sodium, surfactants or emulsifiers and weighting material.

The clays used in drilling fluids containing the base oil compositions of the present invention are suitably organophilic clays. If a hydrophilic clay such as bentonite is available it is necessary to transform these into an organophilic condition e.g. by reaction with appropriate organic ammonium salts by methods well known in the art. These clays act as viscosifiers and the drilling fluid is usually prepared by mixing the conventional components with the base oil composition followed by shearing to achieve the desired rheology, i.e. a fairly low plastic viscosity (PV) and a high yield point (YP) to plastic viscosity ratio, that is YP/PV

During preparation of a drilling fluid, a preformed base oil composition containing both the low toxicity oil component and the polar activator may be mixed with the other conventional components of such fluids such as clay, water, salts, emulsifiers etc. On the other hand the two components of the base oil composition of the present invention may be mixed separately with the other drilling fluid components during agitation or shearing. In the latter case it is preferable to first mix all the polar activator and from one-quarter to one-half of the total low toxicity oil component with the clay and other ingredients. This results in a rapid swelling and gelation of the clay within a few seconds. Thereafter the remainding three quarters of the low toxicity oil component can be added to the swollen gel followed by shearing to achieve a drilling fluid of desired rheology, i.e. a very high YP/PV ratio.

The use of a combination of a low toxicity oil, i.e. an oil component low (less than 10% w/w) in aromatic hydrocarbon content, and a polar activator as the base oil composition can be further enhanced by reducing the amount of emulsifiers used in the drilling fluids. The resultant drilling fluids show a yield point only marginally less than those using diesel as the base oil and yet retain the advantages of low toxicity and low plastic viscosity.

The base oil compositions of the present invention and the use thereof in drilling fluids is further illustrated with reference to the following Examples.

Examples

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In order to demonstrate the relative effectiveness of alcohol ethers as polar activators, other proprietary compounds such as propylene carbonate were also tested for comparison. The results of these tests are tabulated below:-

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TABLE 1

Materials Used

Base Oil	Composition paraffin: P naphthene: N aromatic A	Boiling Range °C	Flash Point (PMCC) °C	Kinematic Viscosity at 40°C cS
Low	P:N:A			
Toxicity	56:42:2	195-255	72	1.7
Kerosine*				
Low	P:N:A			
Toxicity	74:23:3	210-312	109	2.5
Technical	49:42:9	218-310	100	2.9
Gas Oil*				
Diesel**	P:N:A			
- 1	42:37:21	200-365	65-75	3.3

*Some variation in composition and physical properties occur due to variations in crude oil feedstock.

**Sample used was not analysed. Properties shown above are for a typical diesel oil.

TABLE 2

Potar Activators Used 40 **Polar Activators** Approximate Composition Phenoxypropanol (PhP) Phenoxypropoxypropanol (PhDP) Mixed Phenoxypropanol/Phenoxypropoxy Essentially a mixture of PhP/PhDP (95/5 w/w) 45 propanol Essentially IBP 75%, IBDP 21% and others 4%). Mixed Isobutoxypropanol (IBP)/isobutoxypropoxy propanol (IBDP) Phenoxypropanol (PhP) + PhP is the activator and EDP is the cosolvent Ethoxypropoxypropanol (EDP) 50 Acetophenone Propylene carbonate

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TABLE 3

	Cartiful and the state of the s
	materials sources (and meir role in drilling littles)
Other Standard Drilling	Remarks
Fluid Components	
Geltone II*	A gel-forming organophilic clay with bentonitic structure.
Duratone H7*	An oil dispersible lignitic colloid used to control the filtration of oil muds. It also aids in solids
	suspension and emulsification.
Calcium Chloride	This is dissolved in the water phase to provide osmotic force to dehydrate water wet
	formations and increases the mud density.
Lime	Essential to adjust pH and ensure optimum performance of the emulsifiers.
Invermul-L*	Maleic-tall oil amide in kerosine solvent. This is a water-in-oil primary emulsifier.
Ezmul	Partial amide of polyamine and fatty acid in kerosine solvent. It is a secondary emulsifier and
	oil-wetting agent which ensures that baryte and drilled solids do not become water-wetted.
Baryte	Barium sulphate (specific gravity 4.5) is used as weighting agent.

*Registered Trade Mark

1. Rheology Measurement

The characteristics of a drilling fluid can be closely defined by quoting values for PV, YP and gel strength. Standard field and laboratory tests have been devised which are quick and practical. Such tests only approximately reflect downhole behaviour, but they serve their purpose if their limitations are understood and if the data obtained from them are correlated with experience.

A Fann viscometer was used in this study to determine the rheology of low toxicity kerosine, low toxicity technical gas oil and diesel based drilling fluids. See Section 3 below for more details.

2. Standard Drilling Fluid Preparation (General Procedure)

The drilling fluid base oil and polar activator, if any, were measured into the mixing vessel and agitation started using an Ystral mixer. A reduced shear rate (Variac controller setting: 100v) was used for all stages until the final addition of barytes. Geltone and Duratone powders were added over a period of 30 seconds. The mixture was sheared for 10 minutes.

The calcium brine was added and shearing was continued for a further 20 minutes.

The lime, Invermul and Ezmul were added and the shearing was continued for an additional 5 minutes.

The shear rate was increased (Variac setting : 120v) and the baryte was added incrementally over one or two minutes. Finally the complete drilling fluid was sheared at the higher rate for 30 minutes. The vigorous mixing caused the drilling fluid to heat up; the final temperature being 60 ± 5 °C.

When circulating in the oil well, the drilling fluid is exposed to much more severe shearing through the drilling bit at downhole temperatures of 150°C or more. Under these conditions, the organophilic clay will yield to provide maximum structural properties after a few circulations. For specific amounts of various components used see Tables 4 and 6 below.

TABLE 4

30	Drilling Fluids - Their Composition and Pre Compositio	
		Quantities used
_	Lox toxicity oil or diesel	270 ml
35	Polar activator (if any)	5.5 ml (5.8g for PhP; d = 1.06)
	Geltone II	8.0g
	Duratone H7	8.7g
	Calcium brine solution (brine concentration: 463g CaCl ₂ .6H ₂ O diluted to 500 ml H ₂ O)	85 ml
40	Lime	8.0 g
	Invermul solution (Invermul solution conc: 50% vol in base oil) ^b	20 ml
	Ezmul solution (Ezmul solution conc: 50% vol in kerosine)	8.0 ml
45	Baryte	330g
	1 -	1 -

Notes

- a. The drilling fluid composition was varied slightly in some formulations; details of divergences from the above composition are given in Table 6. This formulation is for a 12.2 pounds (weight) per gallon drilling fluid.
- b. Since primary and secondary surfactants (Invermul and Ezmul) were viscous liquids, stock solutions were prepared by diluting to 50% concentration by volume in base oil. The 50% solution was then used in all drilling fluid preparations.

3. Rheology Measurements with the Fann 35SA Direct Indicating Viscometer

This is a concentric cylinder viscometer that enables the variation of shearing stress with shear rate to

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be observed.

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Procedure for Measuring Drilling Fluid Rheology

The freshly prepared drilling fluid was transferred to screw top storage jars. Experience showed that the rheological characteristics of the drilling fluid did not change significantly over a period of 3 months' storage, unless exposed to further heating or shearing.

Just prior to the viscosity measurements, the drilling fluid temperature was dropped 2 or 3 degrees below 20°C. The jar was vigorously shaken to ensure fluid homogeneity. A pre-set volume of drilling fluid was then transferred to the stainless steel beaker of the Fann viscometer. Then the beaker was raised until the drilling fluid surface reached a prescribed mark on the outer cylinder of the viscometer.

Readings were taken at each of six rotation speeds at the moment when the temperature had risen to 20°C, it was found that small divergences from this temperature could significantly affect the viscosity readings.

The drilling fluid was then stirred by switching to maximum rotational speed for 30 seconds. Rotation was stopped for 10 seconds, then re-started at 3 rpm and the maximum momentary reading noted. Rotation was stopped again, but this time for a period of 10 minutes. After this period at rest, the maximum deflection at 3 rpm was noted once more.

Calculation of Results

The following example is for a kerosine drilling fluid with no polar activator (cf Table 6):

TABLE 5

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Viscometer Rotational Speed (rpm) Viscometer Readings at 20 C 600 60 300 33 200 24 100 14 6 3 3 Reading after 10 seconds rest 4 Reading after 10 minutes rest

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In the data tabulated in Table 6 the following abbreviations have been used:

LTK - Low toxicity kerosine

LTGO - Low toxicity technical gas oil

AV - Apparent viscosity (cP)

PV - Plastic viscosity (cP) YP - Yield point (lb/100 ft²)

Unless otherwise indicated as a footnote the standard procedure in Section 2 above was used.

	,		ΔI	rilling Muc	TABLE 6 Drilling Mud Rheologies (20°C)	(20°C)			
¥	Mud Run Number	1	2	3	4	5	9	7	8
Base	Base 011	LTK	LTGO	Diesel	LTK	LTK	LTK	LTK	LTK
Acti	Activator (X weight on base oil)	ł	1		PhP (2.7%)	PhP (2.7%)	PhP (2.7%) PhP (2.7%) PhDP (2.7%) PhDP (2.8%)	PhDP (2.8%)	PhP/PhDP Mixture (2.7%)
Spe	Special Peatures of Mud					(8)		(q)	
	Shear Rate (rpm) 600 300 200 100 6 6 Cel Strength	60 33 24 14 4 4 3	, 85 47 47 34 20 20 6 5 7/12	144 86 66 42 16 14 18/25	74 44 44 44 34 34 23 8 8 7 7	70 44 35 23 9 9 8	75 44 34 23 23 9 8	68 40 31 21 7 6	74 43 34 23 8 8 7 9/11
	Apparent via (AV) (cP)	30	43	7.2	37	35	38	34	37
	Plastic vis (PV) (cP)	27	38	28	93	56	31	28	31
	Yield Point (YP) (1b/100 ft ²)	9	6	58	14	18	13	12	12
	Ratio YP PV	0.22	0.24	0.48	0.47	69.0	0.42	0.43	0.39

Base oil added in two portions such that Geltone initially in contact with more concentrated PhP solution (10%) Same as Run 6 except that slightly more PhDP used and order of addition of PhDP and Geltone to base oil reversed (Geltone added first)

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TABLE 6 - (Continued)
Drilling Mud Rheologies (20°C)

Mud	Mud Run Number	6	10	11	12	13	14	15	16
Вав	Base Oil	LTK	LTK	LTK	רדא	LTK	Diesel	LTGO	LTK
Act	Activator (X weight on base oil)	PhP (5.5%)	•	PhP/PhDP Mixture (2.7%)	PhP/PhDP Mixture (2.7%)	ı	ı	PhDP/PhDP Mixture (2.7%)	IBP/IBDP Mixture (2.7%)
Spe	Special Features of Mud		(c)	(p)	(e)	(f)	(8)	. (H)	
	Shear Rate (rpm)								
	009	77	. 67	75	89	80	126	76	63
	300	44	/ 36	46	27	77	75	28	35
	200	*	56	35	¢3	32	85	97	56
	100	54	16	24	8	19	88	31	17
_	9	6	7	80	11	S	15	11	۰
	م	•	m	9	2	4	14	6	S.
	Gel Strength	8/10	1/8	8/10	12/14	6/11	16/20	10/12	7/9
	Apparent vis (AV) (cP)	39	34	38	57	04	63	47	32
	Plastic vis (PV) (cP)	33	31	29	32	36	51	36	28
	Yield Point (YP) (1b/100 ft ²)	11	'n	17	25	80	24	22	7,
	Ratio YP PV	0.33	0.16	0.59	0.78	0.22	0.47	0.61	0.25

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Drilling Mud Rheologies (20°C) 17 18 19 20 2	35		25 Y B B V F	20	15	10	5
1.5		Dr11	ling Mud Rhe	cologies (20	().		
LTGO LTGO LTGO L 1.5 2.0 2.25 5 PhP PhP PhP PhP PhP PhP PhP PhP PhP PhP	ļ	17	18	19	20	21	22
1.5 2.0 2.25 5 PhP PhP PhP PhP PhP PhP PhP PhP PhP PhP		LTCO	LTGO	LTGO	LTGO	LTGO	LTGO
99 99 105 58 59 62 44 45 47 28 30 31 8 10 9 8 8 8/10 9/11 8/11 50 50 53 41 40 43 17 19 19	1	1.0 PhP	1.5 PhP	2.0 PhP	2.25 PhP	5.4 PhP	2.0 PhP 1.0 EDP
99 99 105 58 59 62 44 45 47 28 30 31 8 10 9 7 8 8 8/10 9/11 8/11 50 50 53 41 40 43 17 19 19							
58 59 62 44 45 47 28 30 31 8 10 9 7 8 8 8 8/10 9/11 8/11 50 50 53 41 40 43 17 19 19		101	66	66	105	95	92
44 45 47 28 30 31 8 10 9 7 8 8 8/10 9/11 8/11 50 50 53 41 40 43 17 19 19 0.41 0.48 0.44		29	28	59	62	26	54
28 30 31 8 10 9 7 8 8 8/10 9/11 8/11 50 50 53 41 40 43 17 19 19		45	77	45	47	42	41
8 10 8 8 8/10 9/11 8/11 50 50 53 41 40 43 17 19 19 0.41 0.48 0.44		53	28	00 ;	31	27	26 8
8/10 9/11 8/11 50 50 53 41 40 43 17 19 19 0.41 0.48 0.44		σ -α	× ^	2 , «	D) 00	٥ ٨	۰ ۸
50 50 53 41 40 43 17 19 19 0.41 0.48 0.44		8/12	8/10	9/11	8/11	8/10	8/10
41 40 43 17 19 19 0.41 0.48 0.44	1	51	20	20	53	87	97
17 19 19 0.41 0.48 0.44		42	41	04	43	39	38
0.41 0.48 0.44		17	17	19	19	17	16
		0.41	0.41	0.48	0.44	0.44	0.42

For Footnotes see page 15.

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5 10		Repeat of standard preparation (see Run 1) except invermul solution halved to 10 ml (see Section 2) Repeat of Run 8, except invermul solution halved to 10 ml Repeat of Run 11 except base oil added in two portions so that Geltone initially in contact with more concentrated polar activator solution (4.4%) Repeat of standard preparation (see Run 1), except Geltone increased by one third (10.7g used instead of 8.0g - see Section 2) See Section 2) See Section 2) Base oil added in two portions so that Geltone initially in contact with more concentrated polar activator (4.4%). Invermul solution halved to 10 ml
15		tun 8, except Invermul solution halved to 10 ml (see Section 2) tun 8, except Invermul solution halved to 10 ml tun 11 except Invermul solution halved to 10 ml tun 11 except Invermul solution (4.4%) tun 11 except base oil added in two portions so that Geltone initially in contact with more candinated preparation (see Run 1), except Geltone increased by one third (10.7g used instead on 2) teses preparation (see Run 3) except Geltone decreased by one third (5.3g used instead of 11 ese 1 two portions so that Geltone initially in contact with more concentrated polar activermul solution halved to 10 ml
20		mul solution ml so that Gelt one increase decreased b y in contact
30		except Inversalved to 10 we portions concept Geltone cone initial
35		(see Run 1) ul solution h 11 added in t olution (4.4% (see Run 1), see Run 3) ex so that Gelt ved to 10 ml
40		f Run 8, except Invermul solution halved to 10 ml f Run 8, except Invermul solution halved to 10 ml f Run 11 except base oil added in two portions so sated polar activator solution (4.4%) tandard preparation (see Run 1), except Gelton fidesel preparation (see Run 3) except Geltone differed in two portions so that Geltone initially Invermul solution halved to 10 ml
45	: .	Repeat of standard preparation (see Run 1) e Repeat of Run 8, except Invermul solution ha Repeat of Run 11 except base oil added in tw concentrated polar activator solution (4.4%) Repeat of standard preparation (see Run 1), - see Section 2) Repeat of diesel preparation (see Run 3) exc See Section 2) Base oil added in two portions so that Gelto (4.4%). Invermul solution halved to 10 ml
50	Footnotes:	(C) Repe (d) Repe (e) Repe (f) Repe ———————————————————————————————————

TABLE 6 - (Concluded) Drilling Mud Rheologies (20°C)

<u>· </u>	DITITING MAD KNEOT	ORIES (20 C	
· Mud	Run Number	23	24
В	ase Oil	LTK	LTK
	tor (% weight on base oil)	Propylene Carbonate (2.7%)	Acetophenone (2.7%)
Special	Features of Mud		
Fann Viscometer Readings	Shear Rate (rpm) 600 300 200 100 6 3 Gel Strength	65 35 25 16 3 2 4/5	68 38 28 18 6 5
Derived	Apparent vis (AV) (cP) Plastic vis (PV)	33 30	34
Rheology Props.	(cP) Yield Point (YP) (1b/100 ft ²)	5	8
	Ratio <u>YP</u> PV	0.17	0.27

Claims

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- 1. A base oil composition suitable for use in drilling fluids, said composition comprising an oil component with an aromatic hydrocarbon content of less than 10% w/w and a polar activator which is an ether alcohol or is a mixture of ether alcohols.
 - 2. A base oil composition according to claim 1 wherein the ether alcohol has one or more ether linkages.
 - A base oil composition according to any one of the preceding claims wherein the ether alcohol is an aliphatic ether alcohol, an aromatic ether alcohol or mixtures thereof.
- 4. A base oil composition according to any one of the preceding claims wherein the ether alcohol is selected from one or more of butoxypropanol, isobutoxypropanol, phenoxypropanol, phenoxypropanol and substituted derivatives thereof whereby the substituents are such that they do not adversely affect the performance of the composition when used in drilling fluids.
 - A base oil composition according to any one of the preceding claims wherein said composition contains from 1-5% w/w of the polar activator.
 - 6. A base oil composition according to any one of the preceding claims wherein the polar activator is a mixture of an aromatic ether alcohol and an aliphatic ether alcohol, the latter being present in the

mixture in an amount from 20-50% w/w of the total mixture.

- 7. A base oil composition according to any one of the preceding claims wherein said composition comprises an oil component, an ether alcohol and one or more of clays, water, halide salts of sodium or calcium, surfactants, emulsifiers and weighting material.
- 8. A drilling fluid comprising the base oil composition as claimed in claim 1 and an organophilic clay.
- 9. A drilling fluid according to claim 8 wherein said fluid is prepared by initially mixing the polar activator and one quarter of the oil component of the base oil composition with the clay, allowing the mixture swell and gel, thereafter adding the remaining three quarters of the oil component to the swollen gel and finally shearing the total mixture to form the drilling fluid.

Revendications

- 15
 - Composition d'huile de base apte à être utilisée dans des fluides de forage, ladite composition comprenant un composant huile ayant une teneur en hydrocarbures aromatiques de moins de 10 % en poids/poids et un activateur polaire qui est un éther-alcool ou qui est un mélange d'éther-alcools.
- Composition d'huile de base suivant la revendication 1, dans laquelle l'éther-alcool porte une ou plusieurs liaisons éther.
 - Composition d'huile de base suivant l'une quelconque des revendications précédentes, dans laquelle l'éther-alcool est un éther-alcool aliphatique, un éther-alcool aromatique ou leurs mélanges.
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- 4. Composition d'huile de base suivant l'une quelconque des revendications précédentes, dans laquelle l'éther-alcool consiste en un ou plusieurs des composés butoxypropanol, isobutoxypropanol, phénoxyéthanol, phénoxypropanol, phénoxypropanol et leurs dérivés substitués dont les substituants sont choisis de manière à ne pas affecter défavorablement la performance de la composition lorsqu'elle est utilisée dans des fluides de forage.
- Composition d'huile de base suivant l'une quelconque des revendications précédentes, qui contient 1 à 5 % en poids/poids de l'activateur polaire.
- 6. Composition d'huile de base suivant l'une quelconque des revendications précédentes, dans laquelle l'activateur polaire est un mélange d'un éther-alcool aromatique et d'un éther-alcool aliphatique, ce dernier étant présent dans le mélange en une quantité de 20 à 50 % en poids par rapport au poids du mélange total.
- 7. Composition d'huile de base suivant l'une quelconque des revendications précédentes, qui comprend un composant huile, un éther-alcool et une ou plusieurs substances telles que des argiles, l'eau, des halogénures de sodium ou de calcium, des surfactants, des émulsionnants et une matière alourdissante.
- 45 8. Fluide de forage comprenant la composition d'huile de base suivant la revendication 1 et une argile organophile.
 - 9. Fluide de forage suivant la revendication 8, qui est préparé par un procédé qui consiste à mélanger initialement l'activateur polaire et un quart du composant huile de la composition d'huile de base avec l'argile, à laisser le mélange gonfler et se gélifier, puis à ajouter les trois quarts restants du composant huile au gel gonflé et finalement à cisailler le mélange total pour former le fluide de forage.

Patentansprüche

Öl-Grundzusammensetzung geeignet zur Verwendung in Bohrflüssigkeiten, wobei die Zusammensetzung eine Ölkomponente mit einem aromatischen Kohlenwasserstoffgehalt von weniger als 10 % Gew./Gew. und einen polaren Aktivator umfaßt, der ein Etheralkohol oder eine Mischung von Etheralkoholen ist.

- 2. Öl-Grundzusammensetzung nach Anspruch 1, worin der Etheralkohol eine oder mehrere Etherbindungen hat.
- 3. Öl-Grundzusammensetzung nach einem der vorstehenden Ansprüche, worin der Etheralkohol ein aliphatischer Etheralkohol, ein aromatischer Etheralkohol oder Mischungen davon ist.
- 4. Öl-Grundzusammensetzung nach einem der vorstehenden Ansprüche, worin der Etheralkohol ausgewählt ist aus einem oder mehreren von Butoxypropanol, Isobutoxypropanol, Phenoxypropanol, Phenoxypropanol und substituierten Derivaten davon, wobei die Substituenten derart sind, daß sie die Leistung der Zusammensetzung nicht nachteilig beeinflussen wenn sie in Bohrflüssigkeiten verwendet werden.
- Öl-Grundzusammensetzung nach einem der vorstehenden Ansprüche, worin die Zusammensetzung von 1 - 5 % Gew/Gew. des polaren Aktivators enthält.
- Öl-Grundzusammensetzung nach einem der vorstehenden Ansprüche, worin der polare Aktivator eine Mischung eines aromatischen Etheralkohols und eines aliphatischen Etheralkohols ist, wobei der letztere in der Mischung in einer Menge von 20 - 50 % Gew./Gew. der gesamten Mischung vorhanden ist.
- 7. Öl-Grundzusammensetzung nach einem der vorstehenden Ansprüche, worin die Zusammensetzung eine Ölkomponente, einen Etheralkohol und ein oder mehrere Tone, Wasser, Halogenidsalze von Natrium oder Calcium, Tenside, Emulgatoren und Beschwerungsmaterial umfaßt.
- 25 8. Bohrflüssigkeit umfassend die Öl-Grundzusammensetzung wie in Anspruch 1 beansprucht und einen organophilen Ton.
- Bohrflüssigkeit nach Anspruch 8, worin die Flüssigkeit hergestellt ist durch erstes Mischen des polaren Aktivators und einem Viertel der Ölkomponente der Öl-Grundzusammensetzung mit dem Ton, Quellenund Gelieren-lassen der Mischung, danach Zusetzen der verbleibenden drei Viertel der Ölkomponente zu dem gequollenen Gel und schließlich Scheren der gesamten Mischung um die Bohrflüssigkeit zu bilden.

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